

## Department of Vermont Health Access Pharmacy Benefits Management Program DUR Board Meeting Agenda

# January 20<sup>th</sup>, 2015 6:00 – 8:30 p.m.

1.	<ul> <li>Executive Session</li> <li>Orientation presentation -GHS</li> <li>Discussion on Medicaid OBRA'90/Supplemental Rebates and A (as provided by 33 VSA § 1998(f)(2))</li> </ul>	<b>6:00 - 6:45</b> greements
2.	Introductions and Approval of DUR Board Minutes (Public Comment Prior to Board Action)	6:45 - 6:45
3.	<b>DVHA Pharmacy Administration Updates</b> <ul><li>Updates</li></ul>	6:45 - 6:50
4.	Medical Director Update	6:50 - 6:55
5.	<b>GHS Presentation on Clinical/Reading Materials</b>	6:55-7:00
6.	Follow-up Items from Previous Meetings <ul><li>Vivitrol step-through for Opiate Dependence</li><li>Zohydro ER new formulation</li></ul>	7:00-7:20
7.	RetroDUR/DUR None this meeting	7:20 – 7:20
8.	Clinical Update: Drug Reviews (Public comment prior to Board action)	7:20 – 7:45
	Abbreviated New Drug Reviews-None	
	<ul> <li>Full New Drug Reviews</li> <li>Entyvio (vedolizumab) 300mg single-dose vial</li> <li>Jublia (efinaconazole) 10% topical solution</li> <li>Sivextro (tedizolid) 200 mg tablet</li> <li>Tanzeum (albiglutide) for injection</li> <li>Harvoni (90 mg ledipasvir and 400 mg sofosbuvir) tablet</li> <li>Viekira pak (ombitasvir, paritaprevir, and ritonavir tablet, dasabe</li> <li>Zohydro ER (hydrocodone bitartrate) extended-release capsules</li> </ul>	uvir tablet)

### 9. Therapeutic Drug Classes – Periodic Review

7:45 - 7:55

(Public comment prior to Board action)
 Sublingual Immunotherapy

#### 10. New Managed Therapeutic Drug Classes

**7:55** – **7:55** 

(Public comment prior to Board action)

None

# 11. Review of Newly-Developed/Revised Clinical Coverage Criteria and/or Preferred Products

7:55 - 8:25

Public comment prior to Board action)

- Hepatitis C update and recommendations
  - Sovaldi
  - Harvoni
  - Viekira Pak

#### 12. General Announcements Selected FDA Safety Alerts

8:25-8:30

- FDA recommends not using lidocaine to treat teething pain and requires new Boxed Warning (6/26/2014 http://www.fda.gov/drugs/drugsafety/ucm402240.htm
- Methylphenidate HCl ER tablets (generic Concerta) made by Mallinckrodt and Kudco: FDA concerns about therapeutic equivalence with 2 generic versions of Concerta tablets (11/13/2014) http://www.fda.gov/drugs/drugs/drugsafety/ucm422568.htm
- FDA reviews long-term antiplatelet therapy as preliminary trial data shows benefits but a higher risk of non-cardiovascular death (11/16/2014) http://www.fda.gov/Drugs/DrugSafety/ucm423079.htm
- FDA warns about case of rare brain infection PML with MS drug Tecfidera (dimethyl fumarate) (11/25/2014) http://www.fda.gov/DrugS/DrugSafety/ucm424625.htm
- FDA issues final rule on changes to pregnancy and lactation labeling information for prescription drug and biological products 12/3/2014)
   <a href="http://www.fda.gov/newsevents/newsroom/pressannouncements/ucm425317.htm">http://www.fda.gov/newsevents/newsroom/pressannouncements/ucm425317.htm</a>
- FDA reporting mental health drug ziprasidone (Geodon) associated with rare but potentially fatal skin reaction (12/11/2014)
   http://www.fda.gov/Drugs/DrugSafety/ucm426391.htm
- o FDA has reviewed possible risks of pain medicine use during pregnancy (1/9/2015) http://www.fda.gov/Drugs/DrugSafety/ucm429117.htm

13. Adjourn 8:30